

the objectives of the Javits-Wagner-O'Day Act (41 U.S.C. 46–48c) in connection with the products proposed for deletion from the Procurement List.

End of Certification

The following products are proposed for deletion from the Procurement List:

Products

Scarf, Branch of Service

NSN: 8455–00–916–8398

NPA: UNKNOWN

Contracting Activity: Defense Supply Center Philadelphia, Philadelphia, PA.

Solvent, Correction Fluid

NSN: 7510–01–333–6241

NPA: Lighthouse for the Blind, St. Louis, MO

Contracting Activity: Federal Acquisition Service, GSA/FSS OFC SUP Ctr—Paper Products, New York, NY.

Patricia Briscoe,

Deputy Director, Business Operations.

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BILLING CODE 6353–01–P

CONSUMER PRODUCT SAFETY COMMISSION

Notice of Meeting of Chronic Hazard Advisory Panel on Phthalates

AGENCY: Consumer Product Safety Commission.

ACTION: Notice of meeting.

SUMMARY: The Consumer Product Safety Commission (Commission) announces the first meeting of the Chronic Hazard Advisory Panel (CHAP) on Phthalates. The Commission appointed this CHAP to study the effects on children's health of all phthalates and phthalate alternatives as used in children's toys and child care articles, pursuant to section 108 of the Consumer Product Safety Improvement Act of 2008 (CPSIA) (Pub. L. 110–314).

DATES: The meeting will be held from 9 a.m. to 5 p.m. on Wednesday, April 14 and from 9 a.m. to 4 p.m. on Thursday, April 15, 2010.

ADDRESSES: The meeting will be held in the fourth floor hearing room in the Commission's offices at 4330 East West Highway, Bethesda, Maryland.

Online Registration: Members of the public who wish to attend the meeting are requested to preregister online at <http://www.cpsc.gov/cgibin/chap.aspx>. This meeting will also be available live via Web cast at <http://www.cpsc.gov/webcast>. Registration is not necessary to view the Web cast.

FOR FURTHER INFORMATION CONTACT:

Michael Babich, Directorate for Health Sciences, Consumer Product Safety

Commission, Bethesda, MD 20814; telephone (301) 504–7253; e-mail mbabich@cpsc.gov.

SUPPLEMENTARY INFORMATION: The Commission has previously investigated potential risks posed to children from phthalate plasticizers, especially di(2-ethylhexyl) phthalate (DEHP) and diisononyl phthalate (DINP), which were used to soften some children's teething, rattles, and toys made from polyvinyl chloride (PVC). Phthalates can leach from such products when they are mouthed by children, causing some phthalates to be ingested. In addition, children and adults can be exposed to phthalates from many sources, including consumer products, food, cosmetics, medical devices, and the environment. Certain phthalates have been shown to cause adverse health effects, including birth defects, in laboratory animals.

Section 108 of the CPSIA permanently prohibits the sale of any “children's toy or child care article” containing more than 0.1 percent of three specified phthalates—DEHP, dibutyl phthalate (DBP), and benzyl butyl phthalate (BBP). Section 108 of the CPSIA also prohibits, on an interim basis, the sale of “toys that can be placed in a child's mouth” or “child care articles” containing more than 0.1 percent of three additional phthalates—DINP, diisodecyl phthalate (DIDP), and dinoctyl phthalate (DNOP).

Moreover, section 108 of the CPSIA requires the Commission to convene a CHAP “to study the effects on children's health of all phthalates and phthalate alternatives as used in children's toys and child care articles.” The CPSIA requires the CHAP to complete an examination of the full range of phthalates that are used in products for children and to: (i) Examine all of the potential health effects (including endocrine disrupting effects) of the full range of phthalates; (ii) consider the potential health effects of each of these phthalates both in isolation and in combination with other phthalates; (iii) examine the likely levels of children's, pregnant women's, and others' exposure to phthalates, based on a reasonable estimation of normal and foreseeable use and abuse of such products; (iv) consider the cumulative effect of total exposure to phthalates, both from children's products and from other sources, such as personal care products; (v) review all relevant data, including the most recent, best-available, peer-reviewed, scientific studies of these phthalates and phthalate alternatives that employ objective data collection practices or employ other objective

methods; (vi) consider the health effects of phthalates not only from ingestion but also as a result of dermal, hand-to-mouth, or other exposure; (vii) consider the level at which there is a reasonable certainty of no harm to children, pregnant women, or other susceptible individuals and their offspring, considering the best available science, and using sufficient safety factors to account for uncertainties regarding exposure and susceptibility of children, pregnant women, and other potentially susceptible individuals; and (viii) consider possible similar health effects of phthalate alternatives used in children's toys and child care articles.

The CHAP's examination must be conducted *de novo*, and the statute contemplates completion of its examination within 18 months of appointment of the CHAP. The CHAP must review prior work on phthalates by the Commission, but the Commission's prior work is not to be considered determinative.

The CHAP must make recommendations to the Commission regarding any phthalates (or combinations of phthalates) in addition to those identified in section 108 of the CPSIA or phthalate alternatives that the panel determines should be declared banned hazardous substances. The Commission selected the CHAP members from scientists nominated by the National Academy of Sciences. See 15 U.S.C. 2077, 2030(b).

The first meeting of the CHAP on Phthalates will be held on April 14 and 15, 2010, in the fourth floor hearing room at the Commission's offices at 4330 East West Highway, Bethesda, Maryland. The meeting will begin at 9 a.m. both days and is scheduled to end at 5 p.m. on April 14 and 4 p.m. on April 15. The meeting is open to the public, space permitting, but no opportunity for public participation in the first meeting is scheduled. There will be an opportunity in connection with the second meeting of the CHAP for presentation of oral and written data and views (date to be announced).

At the first CHAP meeting, the CHAP will choose its Chair and Vice Chair and the CPSC staff will present information on the history of the phthalates project, the scope of the CHAP on phthalates, including a review of the *de novo* examination called for in section 108 (b)(2)(B)(i) through (vii) of the CPSIA, and the CPSC staff's toxicity reviews and other work on phthalates. During the remainder of the meeting, the CHAP will consider how it will proceed and begin its deliberations.

Dated: April 6, 2010.

Todd A. Stevenson,
Secretary, Consumer Product Safety
Commission.

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DEPARTMENT OF DEFENSE

Office of the Secretary

Federal Advisory Committee; Board of Regents of the Uniformed Services University of the Health Sciences

AGENCY: Uniformed Services University of the Health Sciences (USU), DoD.

ACTION: Quarterly meeting notice.

SUMMARY: Under the provisions of the Federal Advisory Committee Act of 1972 (5 U.S.C., Appendix, as amended) and the Sunshine in the Government Act of 1976 (5 U.S.C. 552b, as amended), this notice announces a meeting of the Board of Regents of the Uniformed Services University of the Health Sciences (USU) on May 14, 2010.

DATES: The meeting will be held on Friday, May 14, 2010, from:

9 a.m. to 11:30 a.m. (Open Session).

11:30 a.m. to 12:30 p.m. (Closed Session).

12:30 p.m. to 3:30 p.m. (Open Session).

ADDRESSES: The meeting will be held at the Everett Alvarez Jr. Board of Regents Room (D3001), Uniformed Services University of the Health Sciences, 4301 Jones Bridge Road, Bethesda, Maryland 20814.

FOR FURTHER INFORMATION CONTACT: Janet S. Taylor, Designated Federal Official, 4301 Jones Bridge Road, Bethesda, Maryland 20814; telephone 301-295-3066.

Ms. Taylor can also provide base access procedures.

SUPPLEMENTARY INFORMATION:

Purpose of the Meeting

Meetings of the Board of Regents assure that USU operates in the best traditions of academia. An outside Board is necessary for institutional accreditation.

Agenda

The actions that will take place include the approval of minutes from the Board of Regents Meeting held February 2, 2010; acceptance of reports from working committees; approval of faculty appointments and promotions; and the awarding of post-baccalaureate degrees as follows: Doctor of Medicine, Ph.D. in Nursing Science, Master of Science in Nursing, and master's and

doctoral degrees in the biomedical sciences and public health. The Acting President, USU; the Vice President, USU Office of Research; and the President, Henry M. Jackson Foundation for the Advancement of Military Medicine, will also present reports. These actions are necessary for the University to pursue its mission, which is to provide outstanding health care practitioners and scientists to the uniformed services.

Meeting Accessibility

Pursuant to Federal statute and regulations (5 U.S.C. 552b, as amended, and 41 CFR 102-3.140 through 102-3.165) and the availability of space, most of the meeting is open to the public. Seating is on a first-come basis. The closed portion of this meeting is authorized by 5 U.S.C. 552b(c)(6) as the subject matter involves personal and private observations.

Written Statements

Interested persons may submit a written statement for consideration by the Board of Regents. Individuals submitting a written statement must submit their statement to the Designated Federal Official (*see FOR FURTHER INFORMATION CONTACT*). If such statement is not received at least 10 calendar days prior to the meeting, it may not be provided to or considered by the Board of Regents until its next open meeting. The Designated Federal Official will review all timely submissions with the Board of Regents Chairman and ensure such submissions are provided to Board of Regents Members before the meeting. After reviewing the written comments, submitters may be invited to orally present their issues during the May 2010 meeting or at a future meeting.

Dated: April 5, 2010.

Mitchell S. Bryman,

Alternate OSD Federal Register Liaison Officer, Department of Defense.

[FR Doc. 2010-8052 Filed 4-8-10; 8:45 am]

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DEPARTMENT OF DEFENSE

Department of the Army, Corps of Engineers

Notice of Availability of the Draft Supplemental Environmental Impact Statement for the Proposed Baseload Power Plant, East Kentucky Power Cooperative, Inc., Clark County, KY

AGENCY: U.S. Army Corps of Engineers, DoD.

ACTION: Notice of availability.

SUMMARY: The U.S. Army Corps of Engineers (USACE), in accordance with 42 U.S.C. 4321 to 4370(f), is issuing this notice to advise the public that a Draft Supplemental Environmental Impact Statement (SEIS) has been prepared and is available for review and comment.

DATES: Written comments on the Draft SEIS will be accepted for 45 days following publication of the Environmental Protection Agency's notice of availability for the Draft SEIS in the *Federal Register*. The USACE will hold a public hearing at 7 p.m. (EDT) on June 8, 2010.

ADDRESSES: The Draft SEIS can be viewed online at: <http://www.ekpc.coop/smith-unit1.html>. The Draft SEIS will also be available for viewing at the locations listed in the **SUPPLEMENTARY INFORMATION** section. Comments should be submitted to Mr. Michael Hasty, Acting Chief, South Section, Regulatory Branch, Louisville District, P.O. Box 59, Louisville, KY 40201-0059. The Public Hearing will be held at the Clark County Extension Service office located at 1400 Fortune Drive in Winchester KY 40391.

FOR FURTHER INFORMATION CONTACT: Mr. Michael Hasty, Acting Chief, South Section, Regulatory Branch, Louisville District, P.O. Box 59, Louisville, KY 40201-0059. Phone (502) 315-6676, e-mail: michael.d.hasty@usace.army.mil.

SUPPLEMENTARY INFORMATION: A final EIS and Record of Decision was prepared by the U.S. Department of Energy in 2002/2003 for a proposed 540 megawatt coal-fired integrated gasification combined cycle (IGCC) electric generating plant at the same location, the Smith Site. That project, known as the Kentucky Pioneer IGCC Demonstration Project, was never built. The Corps has reviewed the EIS prepared by DOE and, based on similarities between the two projects, has determined to adopt that EIS as the basis for review of the current proposal. The Corps is preparing this Supplemental EIS (SEIS) to evaluate those aspects of the current proposal that are not substantially similar to the DOE project, as a result of changes in project parameters, existing environmental conditions, and relevant laws and regulations.

East Kentucky Power Cooperative, Inc. (EKPC) has applied for a Department of the Army (DA) permit from the U.S. Army Corps of Engineers (Corps) to authorize unavoidable impacts to jurisdictional waters of the U.S. pursuant to Section 404 of the Clean Water Act (Section 404) and Section 10 of the Rivers and Harbors Act (Section 10).